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APPLICATION NO.	FILI	NG DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/081,457	02/	/21/2002	Anne M. Pianca	98P1021US08	3029
	7590	12/10/2003		EXAMI	NER
PACESETT			EVANISKO, GEORGE ROBERT		
15900 Valley Sylmar, CA				ART UNIT	PAPER NUMBER
•				3762	
				DATE MAILED: 12/10/2003	13

Please find below and/or attached an Office communication concerning this application or proceeding.

-,	Application No.		pplicant(s)	/-					
	Application No			$\cap d$					
	10/081,457	PI	IANCA ET AL.						
Office Action Summary	Examiner	A	rt Unit						
	George R Evani		762						
The MAILING DATE of this communication a Period for Reply	ppears on the cove	r sheet with the corr	espondence add	ress					
A SHORTENED STATUTORY PERIOD FOR REP THE MAILING DATE OF THIS COMMUNICATION - Extensions of time may be available under the provisions of 37 CFR after SIX (6) MONTHS from the mailing date of this communication. - If the period for reply specified above is less than thirty (30) days, a recommendation of the period for reply is specified above, the maximum statutory perions after the period for reply will, by statuses and period for reply will, by statuses and patent term adjustment. See 37 CFR 1.704(b). Status	J. 1.136(a). In no event, how apply within the statutory mi od will apply and will expire ute, cause the application	ever, may a reply be timely to nimum of thirty (30) days will SIX (6) MONTHS from the to to become ABANDONED (3	filed I be considered timely. mailing date of this con U.S.C. § 133).	nmunication.					
1)⊠ Responsive to communication(s) filed on <u>23</u>	September 2003.								
,	is action is non-fin	al.							
3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213.									
Disposition of Claims									
4) Claim(s) 1-17 is/are pending in the application	on.								
4a) Of the above claim(s) is/are withd	4a) Of the above claim(s) is/are withdrawn from consideration.								
5) Claim(s) is/are allowed.	,								
6)⊠ Claim(s) <u>1-17</u> is/are rejected.	6)⊠ Claim(s) <u>1-17</u> is/are rejected.								
7) Claim(s) is/are objected to.									
8) Claim(s) are subject to restriction and	I/or election require	ement.							
Application Papers									
9)☐ The specification is objected to by the Exami	ner.								
10) The drawing(s) filed on is/are: a) □ a	ccepted or b) 🗌 ot	jected to by the Exa	aminer.						
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).									
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).									
11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.									
Priority under 35 U.S.C. §§ 119 and 120									
12)									
Attachment(s)		_							
1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO-1449) Paper No(s)	5)	Interview Summary (PT Notice of Informal Pate Other:							

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DETAILED ACTION

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

Claims 1, 2, 4, 6, 9, 11, 12, and 14 are rejected under 35 U.S.C. 103(a) as being unpatentable over Chastain et al, 5925073, in view of Swoyer, 5683445. Since a guidewire is used in Chastain through the lumen, it is inherent that there be a distal opening in the lead (in the alternative, see the 103 rejection below).

Chastain discloses the claimed invention and providing an anchor in the coronary sinus to stabilize the electrode, but does not teach having a tip electrode and canted portion that orients the tip electrode toward the vessel wall and the number of non-helical bends being exactly two. Swoyer teaches that it is known to have a coronary sinus anchor lead have a tip electrode and canted portion that orients the tip electrode toward the vessel wall to provide effective

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stimulation of the heart. It would have been obvious to one having ordinary skill in the art at the time the invention was made to modify the coronary sinus anchor lead as taught by Chastain, with a tip electrode and canted portion that orients the tip electrode toward the vessel wall as taught by Swoyer, since such a modification would provide a coronary sinus anchor lead with a tip electrode and canted portion that orients the tip electrode toward the vessel wall to provide effective stimulation of the heart.

In addition, it would have been an obvious matter of design choice to a person of ordinary skill in the art to modify the lead as taught by Chastain in view of Swoyer with the lead having exactly two non-helical bends, because Applicant has not disclosed that the lead having exactly two non-helical bends provides an advantage, is used for a particular purpose, or solves a stated problem. One of ordinary skill in the art, furthermore, would have expected Applicant's invention to perform equally well with the lead having a plurality of non-helical bends as taught by Chastain in view of Swoyer, because it provides an effective way to easily anchor the lead.

Therefore, it would have been an obvious matter of design choice to modify Chastain in view of Swoyer to obtain the invention as specified in the claim(s).

Claim 5 is rejected under 35 U.S.C. 103(a) as being unpatentable over Chastain et al in view of Swoyer.

Chastain in view of Swoyer discloses the claimed invention except for the ring electrode located on, before, or after the bends. It would have been obvious to one having ordinary skill in the art at the time the invention was made to modify the anchoring lead as taught by Chastain in view of Swoyer, with the use of a ring electrode on, before or after the bends since it was known

in the art that ring electrodes are included anywhere on leads to provide bipolar sensing and pacing or additional sensing and pacing.

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Claims 10 and 13 are rejected under 35 U.S.C. 103(a) as being unpatentable over Chastain et al in view of Swoyer as applied to claims 6 and 1 above.

Chastain in view of Swoyer discloses the claimed invention except for the humps being in different geometric planes. It would have been an obvious matter of design choice to one skilled in the art to modify the anchoring lead as taught by Chastain in view of Swoyer with the humps in the anchor being located in different geometric planes, since applicant has not disclosed that providing the humps in different geometric planes provides any criticality and/or unexpected results and it appears that the invention would perform equally well with any location of the humps, such as the humps being located in the same plane as taught by Chastain in view of Swoyer to anchor the lead in the coronary sinus.

Claims 3, 7, 8, and 14-17 are rejected under 35 U.S.C. 103(a) as being unpatentable over Chastain in view of Swoyer as applied to claims 2, 6, and 1 above.

Chastain in view of Swoyer discloses the claimed invention except for the lead having a distal opening to receive a guidewire, the stylet having a tapered portion, the first and second bend located in the range of 0.15-0.7 inches from the distal end and first bend, and the lead having a textured region of ePTFE or porous material. It would have been obvious to one having ordinary skill in the art at the time the invention was made to modify the medical electrical lead as taught by Chastain in view of Swoyer with the lead having a distal opening to receive a guidewire, the stylet having a tapered portion, and the lead having a textured region of ePTFE or

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porous material (such as silicone rubber, polyurethane, or ceramic) since it was known in the art that medical electrical leads have a distal opening to receive a guidewire to allow the lead to be positioned in the body, that leads use a stylet with a tapered portion to allow the stylet to fit in the narrow distal end of the lead and to position the lead, and that leads have a textured region of ePTFE or porous material to allow the lead to anchor in the body.

In addition, it would have been an obvious matter of design choice to one skilled in the art to modify the medical electrical lead as taught by Chastain in view of Swoyer to include ePTFE as the textured region and the first and second bends being located 0.15-0.7 inches from the distal end and first bend, since applicant has not disclosed that ePTFE and the first and second bends being located 0.15-0.7 inches from the distal end and first bend provides any criticality and/or unexpected results and it appears that the invention would perform equally well with any biocompatible textured material or any location of the bends, such as silicone rubber, polyurethane or ceramic for allowing the lead to anchor in the body as taught by Chastain in view of Swoyer and in view of one having ordinary skill in the art for allowing the lead to anchor in the coronary sinus or such as the S-shaped or zig-zag shaped lead location of the bends as taught by Chastain in view of Swoyer to allow the lead to anchor in the coronary sinus.

Claims 1, 2, and 4-13 are rejected under 35 U.S.C. 103(a) as being unpatentable over Swoyer in view of Chastain et al (5925073).

Swoyer discloses the claimed invention to anchor a lead in the coronary sinus except for the lead having an s-shape with exactly two non-helical bends for the anchoring and using a conductor to preform the lead into the S-shape. Chastain teaches that it is known to use an s-

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shaped lead with a plurality of non-helical bends to anchor a lead in the coronary sinus and to use the conductor to preform the lead into the S-shape to easily preform the lead using the existing conductor. It would have been obvious to one having ordinary skill in the art at the time the invention was made to modify the lead as taught by Swoyer, with the s-shaped lead with a plurality of bends and preforming the lead with the conductor as taught by Chastain, since such a modification would provide a lead with an s-shape with a plurality of bends to anchor a lead in the coronary sinus and preforming the lead with the conductor into the S-shape to easily configure the lead into a shape using the existing conductor.

In addition, it would have been an obvious matter of design choice to a person of ordinary skill in the art to modify the lead as taught by Swoyer in view of Chastain with the lead having exactly two non-helical bends, because Applicant has not disclosed that the lead having exactly two non-helical bends provides an advantage, is used for a particular purpose, or solves a stated problem. One of ordinary skill in the art, furthermore, would have expected Applicant's invention to perform equally well with lead having a plurality of non-helical bends as taught by Swoyer in view of Chastain, because it provides an effective way to anchor the lead.

Therefore, it would have been an obvious matter of design choice to modify Swoyer in view of Chastain to obtain the invention as specified in the claim(s).

Claims 1, 2, 4, 5, 6, 9, 11 and 12 are rejected under 35 U.S.C. 103(a) as being unpatentable over Alferness et al. Alferness states in column 7, line 60, that his fixation section may also be a "serpentine" configuration (a sinuous or sine curve configuration) and therefore provides non-helical s-shaped bends but does not state he uses exactly 2 bends.

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Alferness discloses the claimed invention except for the lead being shaped to allow the electrode to contact cardiac tissue, using exactly two non-helical bends, and using the conductor to preform the lead. It would have been obvious to one having ordinary skill in the art at the time the invention was made to modify the heart lead as taught by Alferness, with the lead being shaped to allow the electrode to contact cardiac tissue and to preform the lead using the conductor since it was known in the art that heart leads include the lead being shaped to allow the electrode to contact cardiac tissue to effectively stimulate the cardiac tissue and allow selection of a particular area to be stimulated and since it was known in the art that heart leads use the conductor to preform the lead to easily configure the lead into a shape using the existing conductor.

In addition, it would have been an obvious matter of design choice to a person of ordinary skill in the art to modify the lead as taught by Alferness with the lead having exactly two non-helical bends, because Applicant has not disclosed that the lead having exactly two non-helical bends provides an advantage, is used for a particular purpose, or solves a stated problem. One of ordinary skill in the art, furthermore, would have expected Applicant's invention to perform equally well with lead having a plurality of non-helical bends (the serpentine configuration) as taught by Alferness, because it provides an effective way to anchor the lead.

Therefore, it would have been an obvious matter of design choice to modify Alferness to obtain the invention as specified in the claim(s).

Claims 10 and 13 are rejected under 35 U.S.C. 103(a) as being unpatentable over Alferness et al claims 6 and 1 above.

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Alferness discloses the claimed invention except for the humps being in different geometric planes. It would have been an obvious matter of design choice to one skilled in the art to modify the anchoring lead as taught by Alferness with the humps in the anchor being located in different geometric planes, since applicant has not disclosed that providing the humps in different geometric planes provides any criticality and/or unexpected results and it appears that the invention would perform equally well with any location of the humps, such as the humps being located in the same plane as taught by Alferness to anchor the lead in the coronary sinus.

Claims 3, 7, 8, and 14-17 are rejected under 35 U.S.C. 103(a) as being unpatentable over Alferness as applied to claims 2, 6, and 1 above. Alferness discloses the use of a guidewire with the lead and therefore the lead will inherently have a distal opening. In the alternative, see the 103 rejection below.

Alferness discloses the claimed invention except for the lead having a distal opening to receive a guidewire, the stylet having a tapered portion, the first and second bend located in the range of 0.15-0.7 inches from the distal end and first bend, and the lead having a textured region of ePTFE or porous material. It would have been obvious to one having ordinary skill in the art at the time the invention was made to modify the medical electrical lead as taught by Alferness with the lead having a distal opening to receive a guidewire, the stylet having a tapered portion, and the lead having a textured region of ePTFE or porous material (such as silicone rubber, polyurethane, or ceramic) since it was known in the art that medical electrical leads have a distal opening to receive a guidewire to allow the lead to be positioned in the body, that leads use a stylet with a tapered portion to allow the stylet to fit in the narrow distal end of the lead and to

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position the lead, and that leads have a textured region of ePTFE or porous material to allow the lead to anchor in the body.

In addition, it would have been an obvious matter of design choice to one skilled in the art to modify the medical electrical lead as taught by Alferness to include ePTFE as the textured region and the first and second bends being located 0.15-0.7 inches from the distal end and first bend, since applicant has not disclosed that ePTFE and the first and second bends being located 0.15-0.7 inches from the distal end and first bend provides any criticality and/or unexpected results and it appears that the invention would perform equally well with any biocompatible textured material or any location of the bends, such as silicone rubber, polyurethane or ceramic for allowing the lead to anchor in the body as taught by Alferness and in view of one having ordinary skill in the art for allowing the lead to anchor in the coronary sinus or such as the serpentine S-shaped lead location of the bends as taught by Alferness to allow the lead to anchor in the coronary sinus.

Response to Arguments

Applicant's arguments with respect to the claims have been considered but are moot in view of the new ground(s) of rejection necessitated by amendment. The specification does not provide any criticality or unexpected results for using exactly two non-helical bends. The specification teaches away from any criticality of exactly two non-helical bends since "at least two" bends is described and since there are embodiments using three bends.

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Conclusion

Applicant's amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP § 706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to George R Evanisko whose telephone number is 703 308-2612. The examiner can normally be reached on M-F 6:30-5:00.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Angela Sykes can be reached on 703 308-5181. The fax phone number for the organization where this application or proceeding is assigned is 703 306-4520.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is 703 308-1148.

GRE December 8, 2003

GEORGE R. EVANISKO PRIMARY EXAMINER